What is claimed is:

1. A therapeutic composition for the symptomatic relief of cough and nasal congestion associated with adverse respiratory tract conditions in warm-blooded animals in need of such treatment, said composition comprising pharmaceutically effective amounts of active ingredients, wherein said active ingredients consist of phenylephrine tannate, pyrilamine tannate, and guaifenesin.

2. The therapeutic composition of claim 1, in tablet form.

3. The therapeutic composition of claim 2, wherein each tablet contains about 20 to 30 mg of phenylephrine tannate, about 40 to 80 mg of pyrilamine tannate, and about 100 to 400 mg of guaifenesin.

4. The therapeutic composition of claim 2, wherein said tablet form contains about 25 mg of phenylephrine tannate, about 60 mg of pyrilamine tannate, and about 200 mg of guaifenesin.

5. The therapeutic composition of claim 2, wherein said tablet form contains about 25 mg of phenylephrine tannate, about 60 mg of pyrilamine tannate, and about 300 mg of guaifenesin.

6. The therapeutic composition of claim 1, in suspension form.

7. The therapeutic composition of claim 6, wherein said suspension form contains about 3 to 15 mg of phenylephrine tannate, about 25 to 35 mg of pyrilamine tannate, and about 50 to 300 mg of guaifenesin, per 5 ml of suspension.

8. The therapeutic composition of claim 6, wherein said suspension form contains about 5 mg of phenylephrine tannate, about 30 mg of pyrilamine tannate, and about 100 mg of guaifenesin, per 5 ml of suspension.

9. The therapeutic composition of claim 6, wherein said suspension form contains about 5 mg of phenylephrine tannate, about 30 mg of pyrilamine tannate, and about 200 mg of guaifenesin, per 5 ml of suspension.

10. A method for symptomatically treating and relieving the distress of cough and nasal congestion associated with adverse respiratory tract conditions in warm-blooded animals, comprising orally administering to warm-blooded animals in need of such treatment the composition of claim 1.

11. The method of claim 10, wherein said composition is in tablet form.

12. The method of claim 11, wherein each tablet contains about 20 to 30 mg of phenylephrine tannate, about 40 to 80 mg of pyrilamine tannate, and about 100 to 400 mg of guaifenesin.

13. The method of claim 11, wherein said tablet form contains about 25 mg of phenylephrine tannate, about 60 mg of pyrilamine tannate, and about 200 mg of guaifenesin.

14. The method of claim 11, wherein said tablet form contains about 25 mg of phenylephrine tannate, about 60 mg of pyrilamine tannate, and about 300 mg of guaifenesin.

15. The method of claim 10, wherein said composition is in suspension form.

16. The method of claim 15, wherein said suspension form contains about 3 to 15 mg of phenylephrine tannate, about 25 to 35 mg of pyrilamine tannate, and about 50 to 300 mg of guaifenesin, per 5 ml of suspension.

17. The method of claim 15, wherein said suspension contains about 5 mg of

phenylephrine tannate, about 30 mg of pyrilamine tannate, and about 100 mg of guaifenesin, per 5 ml of suspension.

18. The method of claim 15, wherein said suspension contains about 5 mg of phenylephrine tannate, about 30 mg of pyrilamine tannate, and about 200 mg of guaifenesin, per 5 ml of suspension.

19. The method of claim 10, wherein said oral administration is a twice a day administration.